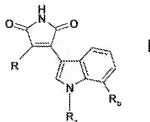


Amendments to the claims.

This listing of claims will replace all prior versions and listings of claims in the application.

Listing of Claims.

1. (currently amended) A compound of formula I



wherein

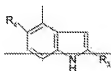
R_a is H; C₁₋₄alkyl; or C₁₋₄alkyl substituted by OH, NH₂, NHC₁₋₄alkyl or N(di-C₁₋₄alkyl)₂;

R_b is H; halogen; C₁₋₆alkyl; or C₁₋₆alkoxy, and

R is a radical of formula (a) or (b)



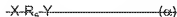
(a)



(b)

wherein

each of R₁ is piperazine and R₂ is a heterocyclic residue; or a radical of formula (c)



wherein X is a direct bond, O, S or NR₁₄, wherein R₁₄ is H or C₁₋₄alkyl;

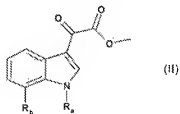
R₆ is C₁₋₄alkylene or C₁₋₄alkylene wherein one CH₂ is replaced by CR_xR_y, wherein one of R_x and R_y is H and the other is CH₃, each of R_x and R_y is CH₃ or R_x and R_y form together CH₂CH₂;

Y is bound to the terminal carbon atom and is selected from OH, NR₁₂R₁₃, wherein each of R₁₂ and R₁₃, independently, is H, C₂₋₆cycloalkyl, C₃₋₆cycloalkyl-C₁₋₄alkyl, aryl, aryl-C₁₋₄alkyl, heteroaryl-C₁₋₄alkyl, C₂₋₆alkenyl or C₁₋₄alkyl optionally substituted on the terminal carbon atom by OH, halogen, C₁₋₄alkoxy or NR₁₄R₁₅, wherein each of R₁₄ and R₁₅, independently, is H, C₁₋₄alkyl, C₃₋₆cycloalkyl, C₃₋₆cycloalkyl-C₁₋₄alkyl, aryl-C₁₋₄alkyl, or R₁₂ and R₁₃ form together with the nitrogen atom to which they are bound a heterocyclic residue; and

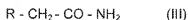
each of R₂ and R₄, independently, is H; halogen; C₁₋₄alkyl; C₁₋₄alkoxy; CF₃; nitrile; nitro or amino,

or a salt thereof.

2. (original) A compound according to claim 1 wherein R_8 is H, methyl, ethyl, or isopropyl, or a salt thereof.
3. (currently amended) A compound according to claim 1 or 2 wherein R_5 is H, Cl, methyl or ethyl, or a salt thereof.
4. (currently amended) A compound according to any one of claims 1 to 3 wherein R_1 is a heterocyclic residue, e.g. a piperazinyl, optionally substituted on a ring nitrogen or on a ring carbon, e.g. 4-methyl-piperazin-1-yl, or 4,7-diaza-spiro[2.5]oct-7-yl; or a radical of formula (II) wherein X is a direct bond, R_6 is CH_3 and Y is $-NR_{12}R_{13}$ wherein each of R_{12} and R_{13} , independently, is H, C_{3-6} cycloalkyl, C_{1-4} alkyl, C_{2-6} alkenyl or C_{1-4} alkyl optionally substituted on the terminal carbon atom by OH, halogen, C_{1-4} alkoxy or $-NR_{14}R_{15}$ wherein each of R_{14} and R_{15} , independently, is H or C_{1-4} alkyl; or R_{12} and R_{13} form together with the nitrogen atom to which they are bound a heterocyclic residue e.g. a piperazinyl, or a salt thereof.
5. (currently amended) A compound according to any one of claims 1 to 4 wherein R_2 and/or R_4 is H; Cl, F_3 , CF_3 ; nitrile; nitro or amino, or a salt thereof.
6. (previously presented): A process for the preparation of a compound of formula I according to claim 1, which process comprises reacting a compound of formula II



wherein R_8 and R_6 are as defined in claim 1,
with a compound of formula III



- wherein R is as defined in claim 1,
and, where required, converting the resulting compound of formula I obtained in free form to a salt form or vice versa, as appropriate.
7. (currently amended) A compound of formula I according to any one of claims 1 to 5, in free form or in a pharmaceutically acceptable salt form for use as a pharmaceutical.
 8. (currently amended) A pharmaceutical composition comprising a compound of formula I according to any one of claims 1 to 5, in free form or in a pharmaceutically acceptable salt form, in association with a pharmaceutically acceptable diluent or carrier therefor.
 9. (cancelled) .

- 10 (currently amended): A method for preventing or treating acute or chronic transplant rejection disorders or diseases mediated by T-lymphocytes and/or PKC- ϵ or GSK-3 β in a subject in need of such treatment, which method comprises administering to said subject an effective amount of a compound of formula I according to any one of claims 1 to 5 or a pharmaceutically acceptable salt thereof.
- 11 (new) A method for preventing or treating T-cell mediated inflammatory or autoimmune diseases, in a subject in need of such treatment, which method comprises administering to said subject an effective amount of a compound of formula I of claim 1 or a pharmaceutically acceptable salt thereof.